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Cys	His	Gln	Leu	Cys	Ala	Arg	Gly	His	Cys	Trp	Gly	Pro	Gly	Pro	
1				5					10					15	
Thr	Gln	Cys	Val	Asn	Cys	Ser	Gln	Phe	Leu	Arg	Gly	Gln	Glu	Cys	
				20					25					30	
Val	Glu	Glu	Cys	Arg	Val	Leu	Gln	Gly	Leu	Pro	Arg	Glu	Tyr	Val	
				35					40					45	
Asn	Ala	Arg	His	Cys	Leu	Pro	Cys	His	Pro	Glu	Cys	Gln	Pro	Gln	
				50					55					60	
Asn	Gly	Ser	Val	Thr	Cys	Phe	Gly	Pro	Glu	Ala	Asp	Gln	Cys	Val	
				65					70					75	
Ala	Cys	Ala	His	Tyr	Lys	Asp	Pro	Pro	Phe	Cys	Val	Ala	Arg	Cys	
				80					85					90	
Pro	Ser	Gly	Val	Lys	Pro	Asp	Leu	Ser	Tyr	Met	Pro	Ile	Trp	Lys	
				95					100					105	
Phe	Pro	Asp	Glu	Glu	Gly	Ala	Cys	Gln	Pro	Cys	Pro	Ile	Asn	Cys	
				110					115					120	
Thr	His	Ser	Cys	Val	Asp	Leu	Asp	Asp	Lys	Gly	Cys	Pro	Ala	Glu	
				125					130					135	
Gln	Arg	Ala	Ser	Pro	Leu	Thr									
				140											

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What is claimed is:

1. A method for extending time to disease progression (TTP) or survival in cancer patients with metastatic breast cancer which displays HER activation comprising: (a) administering a HER2 antibody which comprises the variable light and variable heavy amino acid sequences in SEQ ID Nos. 3 and 4 as fixed doses of 420 mg to the patients and (b) measuring TTP or survival in the patients to confirm it is extended.

2. The method of claim 1 wherein the HER2 antibody inhibits HER heterodimerization.

3. The method of claim 1 wherein the cancer displays HER2 activation.

4. The method of claim 3 wherein the cancer displays HER2 overexpression or amplification.

5. The method of claim 1 wherein the HER2 antibody is pertuzumab.

6. The method of claim 1 wherein the HER2 antibody is a naked antibody.

7. The method of claim 1 wherein the HER2 antibody is an intact antibody.

8. The method of claim 1 wherein the HER2 antibody is an antibody fragment comprising an antigen binding region.

9. The method of claim 1 wherein the HER2 antibody is administered as a single anti-tumor agent.

10. The method of claim 1 comprising administering a second therapeutic agent to the patients.

11. The method claim 10 wherein the second therapeutic agent is selected from the group consisting of chemotherapeutic agent, HER antibody, antibody directed against a tumor associated antigen, anti-hormonal compound, cardioprotectant, cytokine, EGFR-targeted drug, anti-angiogenic agent, tyrosine kinase inhibitor, COX inhibitor, non-steroidal anti-inflammatory drug, farnesyl transferase inhibitor, antibody that binds oncofetal protein CA 125, HER2 vaccine, HER targeting therapy, Raf or ras inhibitor, liposomal doxorubicin, topotecan, taxane, dual tyrosine kinase inhibitor, TLK286, EMD-7200, a medicament that treats nausea, a medicament that prevents or treats skin rash or standard acne

therapy, a medicament that treats or prevents diarrhea, a body temperature-reducing medicament, and a hematopoietic growth factor.

12. The method of claim 10 wherein the second therapeutic agent comprises trastuzumab.

13. The method of claim 1 wherein TTP is extended.

14. The method of claim 1 wherein survival is extended.

15. The method of claim 1 wherein administration of the HER2 antibody extends TTP or survival at least about 20% more than TTP or survival achieved by administering an approved anti-tumor agent to the cancer patients.

16. A method for extending time to disease progression (TTP) or survival in breast cancer patients comprising: (a) administering a HER2 antibody to the patients as fixed doses of about 420 mg of the HER2 antibody so as to extend TTP or survival in the patients, wherein the HER2 antibody comprises the variable light and variable heavy amino acid sequences in SEQ ID Nos. 3 and 4, respectively, and (b) measuring TTP or survival in the patients to confirm it is extended.

17. The method of claim 16 which extends TTP.

18. The method of claim 16 which extends survival.

19. The method of claim 16 wherein the patients are metastatic breast cancer patients.

20. The method of claim 19 wherein the cancer displays HER2 activation.

21. The method of claim 20 wherein the cancer displays amplified or overexpressed HER2.

22. The method of claim 16 wherein the fixed doses of about 420 mg of the HER2 antibody are administered about every three weeks.

23. The method of claim 22 wherein a loading dose of 840 mg of the HER2 antibody is administered followed by the fixed doses of about 420 mg of the HER2 antibody.

24. The method of claim 16 comprising administering a second therapeutic agent to the patients.

25. The method of claim 24 wherein the second therapeutic agent comprises trastuzumab.